PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (Chapter I of the Patent Cooperation Treaty)

(PCT Rule 44bis)

Applicant's or agent's file reference SRI-008PC	FOR FURTHER ACTION	See item 4 below	
International application No. PCT/US2005/002033	International filing date (day/month/year) 21 January 2005 (21.01.2005)	Priority date (day/month/year) 23 January 2004 (23.01.2004)	
International Patent Classification (8th edition unless older edition indicated) See relevant information in Form PCT/ISA/237			
Applicant SRI INTERNATIONAL			

1.	This international preliminary report on patentability (Chapter I) is issued by the International Bureau on behalf of the International Searching Authority under Rule 44 bis. 1(a).				
2.	This REPORT consists of a total of 6 sheets, including this cover sheet. In the attached sheets, any reference to the written opinion of the International Searching Authority should be read as a reference to the international preliminary report on patentability (Chapter I) instead.				
3.	This report contains indications	relating to the following item	is:		
	Box No. I	Basis of the report			
	Box No. II	Priority			
	Box No. III	Non-establishment of opin	nion with regard to novelty, inventive step and industrial		
	Box No. IV	Lack of unity of invention	1		
	Box No. V	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			
	Box No. VI	Certain documents cited			
	Box No. VII	Certain defects in the inte	rnational application		
	Box No. VIII	Certain observations on th	e international application		
4.	4. The International Bureau will communicate this report to designated Offices in accordance with Rules 44bis.3(c) and 93bis.1 but not, except where the applicant makes an express request under Article 23(2), before the expiration of 30 months from the priority date (Rule 44bis .2).				
	-				
			Date of issuance of this report 24 July 2006 (24.07.2006)		
	The International Burn 34, chemin des Co 1211 Geneva 20, S	lombettes	Authorized officer Yolaine Cussac		
Facsin	acsimile No. +41 22 338 82 70 e-mail: pt11@wipo.int				

Form PCT/IB/373 (January 2004)

PATENT COOPERATION TREATY

From the INTERNATIONAL SEAR To:	CHING AUTH	ORITY	1	7.	REC'D 1	4 JUL 2005
MICHAEL A. RODRIGUEZ GUERIN & RODRIGUEZ, LLP		PCWPO POT				
5 MOUNT ROYAL AVE MOUNT ROYAL OFFICE	NUE		w	RITTEN OPI	NION OF	THE
MARLBOROUGH, MA 01752					AUTHORITY	
				(PCT Rule	43 <i>bis</i> .1)	
·			Date of mailing (day/month/year)	12 JUL	2005	
Applicant's or agent's file	reference		FOR FURTHER	ACTION		,
SRI-008PC				See paragraph 2	2 below	
International application N	lo.	International filing date	(day/month/year)	Priority date (day/month/	year)
PCT/US05/02033 International Patent Classi	fication (IPC)	21 January 2005 (21.01.	2005)	23 January 20	04 (23.01.2	004)
IPC(7): B01L 3/00, 3/02; (Applicant	JUIN 21/29, 23	722, 21700, 1710 and US (21.: 422/99, 100, 82.0	5, 930; 436/157,	164, 180	
SRI INTERNATIONAL						
1. This opinion contains	indications rela	ting to the following items	s:			
Box No. I	Basis of the	opinion				
Box No. II	Priority					
Box No. III	Non-establis	hment of opinion with reg	ard to novelty inven	tive step and ind	luct ri ol on al	inchille.
Box No. IV		y of invention	,	ave step and me	usurar appi	icability
Box No. V	Reasoned sta	tement under Rule 43bis. citations and explanation	l(a)(i) with regard to s supporting such sta	novelty, inventi	ve step or i	ndustrial
Box No. VI	Certain docu		,,			
Box No. VII	Certain defec	ts in the international app	lication			
Box No. VIII	Certain obser	vations on the internation	al application			
2. FURTHER ACTIO	N					
Authority other than th	is one to be th	nary examination is made Authority ("IPEA") exc e IPEA and the chosen IF nal Searching Authority w	ept that this does in	not apply where		
ILDIE & WILLOW TOPIN TO	or before the ex	considered to be a written appropriate, with amendme piration of 22 months from /220.	enis before the evni	ration of 2 mont	ha fram the	to submit to the date of mailing
3. For further details, see n						
Name and mailing address of	the ISA/ US		Authorized officer	1		1
Mail Stop PCT, Attn: Commissioner for Pat	ISA/US		Brian Gordon	Mint	11/1	1/9/
P.O. Box 1450 Alexandria, Virginia 2	22313-1450			my.	VVU	1
rm PCT/ISA/237 (cover she	ס		Telephone No. (571) 272-1700		

International application No.	
PCT/US05/02033	

Box No	o. I Basis of this opinion
1. With a was fi	regard to the language, this opinion has been established on the basis of the international application in the language in which i led, unless otherwise indicated under this item.
Ш	This opinion has been established on the basis of a translation from the original language into the following language, which is the language of a translation furnished for the purposes of international search (under Rules 12.3 and 23.1(b)).
2. With r	regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed ion, this opinion has been established on the basis of:
a.	type of material
	a sequence listing
	table(s) related to the sequence listing
b.	format of material
	in written format
	in computer readable form
c.	time of filing/furnishing
	contained in international application as filed.
	filed together with the international application in computer readable form.
	furnished subsequently to this Authority for the purposes of search.
	In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
. Additio	nal comments:
- DCTG	SA/237(Box No. 1) (January 2004)

International application No. PCT/US05/02033

. Statement		
Novelty (N)	Claims 4-9, 11, 13-21, 23-28, 30, 32-36	•
, ,	Claims 1-3, 10, 12, 22, 29, 31	Y N
Inventive step (IS)	Claims 4-9,11, 13-21, 23-28, 30, and 32-36	Y
	Claims 1-3, 10, 12, 22, 29, 31	N
Industrial applicability (IA)	Claims 1-36	Y
	Claims NONE	N
Citations and explanations:		
ease See Continuation Sheet		
•		
	•	
,		

International application No. PCT/US05/02033

Supplemental Box In case the space in any of the preceding boxes is not sufficient.	

V. 2. Citations and Explanations:

Claim 22 and 29 lacks novelty under PCT Article 33(2) as being anticipated by Sasaki et al. US 2003/0086824.

Sasaki et al. diclsoses when a plate for the flat cell transmits measuring light to a certain extent, measurement is carried out by the use of the transmittance or the absorbance calculated on the basis of the transmittance. Specifically, measuring light 8 emitted by a light source 7 passes through the droplet on the hydrophilic pattern and the plate and a portion thereof is absorbed by them. The unabsorbed light 9 reaches a light receptor 10. A specific component in the sample is quantitated from the attenuation factor of the measuring light at a measuring wavelength. When a plate for the flat cell reflects measuring light to a certain extent, measurement is carried out by the use of the reflectance. Specifically, measuring light 12 emitted by a light source 11 passes through the droplet on the hydrophilic pattern, is reflected from the cell surface and passes through the droplet (droplet on surface) on the hydrophilic pattern again. In this process, a portion of the measuring light is absorbed by the droplet and the cell surface. The unabsorbed light, i.e., light 13 that has been reflected from the cell surface and has again passed through the droplet on the hydrophilic pattern reaches a light receptor 14. A specific component in the sample is quantitated from the attenuation factor of the measuring light at a measuring wavelength. See [0045]

Claim 22 and 29 lacks novelty under PCT Article 33(2) as being anticipated by Hess et al. US 2002/0001544 A1.

Hess et al. dicloses at least one operation may be performed on each droplet from the group of operations consisting of mixing, diluting, concentrating, filtering, and analyzing. Analyzing may include performing at least one operation from the group of operations consisting of optical interrogation and mass spectrometry. Optical interrogation may include at least one of fluorescence spectrometry, Raman spectroscopy and UV absorption. Analyzing the content of each droplet may include aspirating each droplet into a dispensing unit and presenting each droplet for analysis via the dispensing unit. Each droplet may be presented to a mass spectrometer and a characteristic of each droplet determined by means of mass spectrometry. Analyzing a characteristic of each droplet may include heating each droplet, or applying a pneumatic or explosive force to each droplet, so as to form an atomized spray and determining a characteristic each droplet by means of mass spectrometry. Each droplet may be vibrated so as to cause atomization, whereupon a characteristic of each droplet can be determined by means of mass spectrometry. Vibrating the droplet may include focusing a pulsed laser (light source) onto the surface or backside of the surface in a proximity of each droplet, utilizing acoustic waves, or mechanically vibrating the surface. A voltage to the surface onto which each droplet is deposited may be applied to assist in the formation of atomized spray. See [0009]

Claim 22 and 29 novelty under PCT Article 33(2) as being anticipated by Anderson et al. US 6,620,620.

Anderson et al. disclose in FIGS. 2A and 2B illustrate the arrangements used in the droplet sensing method known as video drop sensing. Droplet 42 is deposited from needle 36 of syringe 34 onto deposition surface plate 44. Light source 66 directs light through light diffuser 68, droplet 42 and optional diffuser 70 to video camera 72. As shown in FIG. 2B, video screen 74 (which may be a component of display 24) includes crosshairs 76 which are movable by controls 26 so as to permit the operator to determine a change in the light obscuration as the droplet evaporates. Video camera 72 is interfaced with the power, logic and computing module 22 shown in FIG. 1 so that a change in light obscuration due to the evaporation of at least a portion of the liquid or solvent permits the next successive droplet to be deposited in a timely manner. In a preferred embodiment, a blue or other colored filter may be used to improve the sensitivity of the video camera, and a blunt end needle may be used to improve the wetting of the droplet to the needle.

International application No. PCT/US05/02033

Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

FIG. 3 is a visual representation of the sensing of the evaporative process by means of the change in a reflected laser beam. Droplet 42 is deposited from needle 36 of syringe 34 onto deposition surface plate 44. Laser 78 directs a laser beam at the droplet on deposition surface plate 44, which beam is reflected toward detector 80. When the liquid or solvent droplet has not yet fully evaporated, the reflected beam is scattered and the detector 80 does not yet receive the full beam. Detector 80 is interfaced with the power, logic, and computing module 22 shown in FIG. 1 so that a change in the spectral distribution of the reflected laser beam due to the evaporation of at least a portion of the liquid or solvent permits the next successive droplet to be deposited in a timely manner.

Claims I, 3, 10, 12, 22, 29 and 31 novelty under PCT Article 33(2) as being anticipated by Yuguchi et al. US 5,275,787.

Yuguchi et al. discloses In FIGS. 9 and 10, a fixed transparent glass plate 16 is arranged tilted relative to the discharge axis 1 of the nozzle 1. The laser beam emitted from the laser light source 7 is incident from the back of the glass plate 16. The discharge axis 1 of the nozzle 1 and the optical axis O of the laser light are arranged so as to cross near the glass plate 16. The detector 5 is arranged on the optical axis of the laser light. A condenser lens (not shown) and a beam stopper (not shown) for preventing the laser beam from directly entering the detector 5 are provided in front of the detector 5 on the optical axis. The condenser lens and beam stopper form a dark-field optical system so that light scattered in forward directions of the optical axis by a particle S situated at a measuring position on the glass plate 16 onto which the laser beam is projected is subjected to photometry by the detector 5. The detector 6 is arranged in a direction crossing each of the optical axis O of the laser light and the axis 1 of the liquid drop discharge. A condenser lens (not shown) and the wavelength selection filter 15 are provided in front of the detector 6 so that fluorescence emitted from the particle S at the measuring position is selectively subjected to photometry by the detector 6.

The size of the opening and the capacity of the heater are set so that the liquid drops discharged from the nozzle have diameters of about $50 \mu m$ -80 μm .

Claims 22 and 29 novelty under PCT Article 33(2) as being anticipated by Krause et al. US 5,586,200.

Krause et al. teach a system for measuring sample volumes of droplets using a rod-like transfer element. The light of lamp (1) is split into two beams (2) and (3) with each beam being directed onto the sample adhering to the transfer element (4) via a system of lenses, diaphragms, and mirrors. The axis of a rod-like transfer element runs perpendicularly to the drawing surface. The light beams (2) and (3) consequently illuminate the sample droplets from two perpendicular directions which in turn run perpendicularly to the axis of the transfer element. The illuminated sample droplet is located in the focus of the lens systems (5) and (6) by which it is imaged in such a way that a sharp picture is generated on the CCD-camera (8). The bundles of beams emerging from the lens systems (5) and (6) pass through an arrangement of mirrors, diaphragms, and lenses. The semi-permeable mirror (7) directs the partial beams onto the CCD-camera (8).

Claims 4-9, 11, 13-21, 23-28, 30, and 32-36 meet the criteria set out in PCT Article 33(2)-(3), because the prior art does not teach or fairly suggest and immiscible organic liquid that controls evportation, a second immiscible liquid, a thermal gradient in the immiscible liquid, and a dye droplet in the immiscible liquid.

Claims 1-36 meet the criteria set out in PCT Article 33(4), and thus meet industrial applicability because the subject matter claimed can be made or used in industry.

PATENT COOPERATION TREATY

To: MICHAEL A. RODRIGUEZ GUERIN & RODRIGUEZ, LLP 5 MOUNT ROYAL AVENUE		PC WPO PST		
MOUNT ROYAL AVENUE MOUNT ROYAL OFFICE PARK MARLBOROUGH, MA 01752		WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY		
				(PCT Rule 43bis.1)
			Date of mailing (day/month/year)	. 12 JUL 2005
Applicant's or agent's file SRI-008PC	reference		FOR FURTHER	ACTION See paragraph 2 below
International application ?	No.	International filing date	(day/month/year)	Priority date (day/month/year)
PCT/US05/02033	·	21 January 2005 (21.01.	2005)	23 January 2004 (23.01.2004)
		or both national classificat		
(PC(7): B01L 3/00, 3/02; Applicant	G01N 21/29, 2	5/22, 21/00, 1/10 and US (Cl.: 422/99, 100, 82.0	95, 930; 436/157, 164, 180
SRI INTERNATIONAL				
IN INTERNATIONAL				
1. This opinion contains	indications rela	ating to the following item	s:	
Box No. [Basis of the	opinion		
Box No. II	Priority	•		
Box No. III	Non-establis	shment of opinion with rec	rard to novelty inven	tive step and industrial applicability
Box No. IV		y of invention	and to noverty, mivel	mve step and industrial applicability
		J		
Box No. V	Reasoned st	atement under Rule 43 <i>hi</i> e	1/aVi) with record to	manufacture of the second
Box No. V	Reasoned stapplicability	atement under Rule 43bis.; citations and explanation	l(a)(i) with regard to s supporting such sta	novelty, inventive step or industrial
Box No. V	Reasoned strapplicability Certain docu	; citations and explanation	l(a)(i) with regard to as supporting such sta	novelty, inventive step or industrial tement
	applicability Certain docu	; citations and explanation	s supporting such sta	novelty, inventive step or industrial stement
Box No. VI	applicability Certain docu Certain defe	; citations and explanation iments cited	is supporting such sta	novelty, inventive step or industrial atement
Box No. VI Box No. VII	applicability Certain docu Certain defe Certain obse	; citations and explanation iments cited cts in the international app	is supporting such sta	novelty, inventive step or industrial stement
Box No. VI Box No. VII Box No. VIII Box No. VIII 2. FURTHER ACTIO If a demand for international Prelimina Authority other than the	applicability Certain docu Certain defe Certain obse N ational prelimitary Examining sis one to be the	r; citations and explanation rments cited cts in the international app rvations on the internation nary examination is made	lication al application e, this opinion will be that this does a per that the does a	be considered to be a written opinion of the not apply where the applicant chooses an
Box No. VI Box No. VII Box No. VIII Box No. VIII 2. FURTHER ACTIO If a demand for intern International Prelimina Authority other than the that written opinions of If this opinion is, as proposed in the pro	applicability Certain docu Certain defe Certain obse N ational prelimi ary Examining ary Examining for this Internatio rovided above, ogether, where or before the ex	ry citations and explanation aments cited cts in the international approvations on the internation mary examination is made; Authority ("IPEA") except IPEA and the chosen II nal Searching Authority where the internation is made; Authority which is a written appropriate, with amendments of 22 months from	dication al application the this opinion will be that this does in the period of the	the considered to be a written opinion of the not apply where the applicant chooses an International Bureau under Rule 66.1bis(b) ed.
Box No. VI Box No. VII Box No. VIII Box No. VIII 2. FURTHER ACTIO If a demand for intern International Prelimina Authority other than the that written opinions of If this opinion is, as pi IPEA a written reply to	applicability Certain docu Certain defe Certain obse N ational prelimi ary Examining ary Examining for this Internatio rovided above, ogether, where or before the ex	ry citations and explanation aments cited cts in the international approvations on the internation mary examination is made; Authority ("IPEA") except IPEA and the chosen II nal Searching Authority where the internation is made; Authority which is a written appropriate, with amendments of 22 months from	dication al application the this opinion will be that this does in the period of the	the considered to be a written opinion of the not apply where the applicant chooses an International Bureau under Rule 66.1bis(b) ed.
Box No. VI Box No. VII Box No. VIII Box No. VIII 2. FURTHER ACTIO If a demand for international Prelimina Authority other than that written opinions of If this opinion is, as proposed in the property of Form PCT/ISA/220 of For further options, see	Certain docu Certain defe Certain obse N actional prelimitary Examining his one to be the first Internation revided above, begether, where or before the experimental present the experimental pres	ry citations and explanation aments cited cts in the international approvations on the internation mary examination is made (Authority ("IPEA") excite IPEA and the chosen II and Searching Authority we considered to be a writte appropriate, with amendment of 22 months from 1/220.	dication al application the this opinion will be that this does in the period of the	the considered to be a written opinion of the not apply where the applicant chooses an International Bureau under Rule 66.1bis(b) ed.
Box No. VI Box No. VII Box No. VIII 2. FURTHER ACTIO If a demand for intern International Prelimina Authority other than that written opinions of If this opinion is, as proposed in the p	applicability Certain docu Certain defe Certain obse N ational prelimi ary Examining ary Examining is one to be the fithis Internatio rovided above, ogether, where or before the ex Form PCT/ISA notes to Form P	ry citations and explanation aments cited cts in the international approvations on the internation mary examination is made (Authority ("IPEA") excite IPEA and the chosen II and Searching Authority we considered to be a writte appropriate, with amendment of 22 months from 1/220.	dication al application the this opinion will be that this does in the period of the	the considered to be a written opinion of the not apply where the applicant chooses an International Bureau under Rule 66.1bis(b) ed.
Box No. VI Box No. VII Box No. VIII Box No. VIII 2. FURTHER ACTIO If a demand for intern International Prelimina Authority other than the that written opinions of If this opinion is, as proposed in the	applicability Certain docu Certain defe Certain obse ON actional prelimitary Examining his one to be the order of this Internation rovided above, ogether, where or before the examiner of the Examinity for the ISA/US EISA/US	ry citations and explanation aments cited cts in the international approvations on the internation mary examination is made (Authority ("IPEA") excite IPEA and the chosen II and Searching Authority we considered to be a writte appropriate, with amendment of 22 months from 1/220.	dication al application al application be, this opinion will be that this does a PEA has notified the fill not be so consider an opinion of the IPE tents, before the expinant the priority date, we have a consider and the priority date.	the considered to be a written opinion of the not apply where the applicant chooses an International Bureau under Rule 66.1bis(b) ed.
Box No. VI Box No. VII Box No. VIII 2. FURTHER ACTIO If a demand for intern International Prelimina Authority other than that written opinions of If this opinion is, as proposed in the p	applicability Certain docu Certain defe Certain obse N ational prelimi ary Examining his one to be the first internatio revided above, hogether, where hor before the ex Form PCT/ISA motes to Form P	ry citations and explanation aments cited cts in the international approvations on the internation mary examination is made (Authority ("IPEA") excite IPEA and the chosen II and Searching Authority we considered to be a writte appropriate, with amendment of 22 months from 1/220.	lication al application e, this opinion will be that this does open that this does open that the does open that the solution of the IPE tents, before the expirit on the priority date, when the priority date, when the supplication of the IPE tents, before the expirit of the priority date, when the priority date, when the supplication of the IPE tents, before the expirit of the priority date, when the priority date, when the supplication is supplied to the su	the considered to be a written opinion of the not apply where the applicant chooses an International Bureau under Rule 66.1bis(b) ed.

International application No.

PCT/US05/02033

BOXIV	o. I Basis of this opinion
1. With was f	regard to the language, this opinion has been established on the basis of the international application in the language in which it iled, unless otherwise indicated under this item.
	This opinion has been established on the basis of a translation from the original language into the following language which is the language of a translation furnished for the purposes of international search (under Rules 12.3 and 23.1(b)).
2. With inven	regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed tion, this opinion has been established on the basis of:
a.	type of material
	a sequence listing
	table(s) related to the sequence listing
b.	format of material
	in written format
	in computer readable form
. c.	time of filing/furnishing
	contained in international application as filed.
	filed together with the international application in computer readable form.
	furnished subsequently to this Authority for the purposes of search.
3.	In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
4. Additi	oñal comments:
•	·
m PCT/I	SA/237(Box No. 1) (January 2004)

Form PCT/ISA/237 (Box No. V) (January 2004)

International application No. PCT/US05/02033

Box No. V Reasoned statement under Rule 43 bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
1. Statement				
Novelty (N)	Claims 4-9, 11, 13-21, 23-28, 30, 32-36	YES		
	Claims 1-3, 10, 12, 22, 29, 31	NO		
Inventive step (IS)	Claims 4-9,11, 13-21, 23-28, 30, and 32-36	VEC		
	Claims 1-3, 10, 12, 22, 29, 31	YES NO		
Industrial applicability (IA)	Claims 1-36	YES		
	Claims NONE	NO		
Citations and explanations:				
lease See Continuation Sheet				
•				
		•		
		•		
	•			
	•			
•				

International application No. PCT/US05/02033

1	Supplemental Box
ŀ	In case the space in any of the preceding boxes is not sufficient.
l	
l	
ĺ	
	V. 2. Citations and Explanations:
	Claim 22 and 29 lacks novelty under PCT Article 33(2) as being anticipated by Sasaki et al. US 2003/0086824.
	Sasaki et al. diclsoses when a plate for the flat cell transmits measuring light to a certain extent, measurement is carried out by the use of the transmittance or the absorbance calculated on the basis of the transmittance. Specifically, measuring light 8 emitted by a light source 7 passes through the droplet on the hydrophilic pattern and the plate and a portion thereof is absorbed by them. The unabsorbed light 9 reaches a light receptor 10. A specific component in the sample is quantitated from the attenuation factor of the measuring light at a measuring wavelength. When a plate for the flat cell reflects measuring light to a certain extent, measurement is carried out by the use of the reflectance. Specifically, measuring light 12 emitted by a light source 11 passes through the droplet on the hydrophilic pattern, is reflected from the cell surface and passes through the droplet (droplet on surface) on the hydrophilic pattern again. In this process, a portion of the measuring light is absorbed by the droplet and the cell surface. The unabsorbed light, i.e., light 13 that has been reflected from the cell surface and has again passed through the droplet on the hydrophilic pattern reaches a light receptor 14. A specific component in the sample is quantitated from the attenuation factor of the measuring light at a measuring wavelength. See [0045]
	Claim 22 and 29 lacks novelty under PCT Article 33(2) as being anticipated by Hess et al. US 2002/0001544 A1.
	Hess et al. dicloses at least one operation may be performed on each droplet from the group of operations consisting of mixing, diluting, concentrating, filtering, and analyzing. Analyzing may include performing at least one operation from the group of operations consisting of optical interrogation and mass spectrometry. Optical interrogation may include at least one of fluorescence spectrometry, Raman spectroscopy and UV absorption. Analyzing the content of each droplet may include aspirating each droplet into a dispensing unit and presenting each droplet for analysis via the dispensing unit. Each droplet may be presented to a mass spectrometer and a characteristic of each droplet determined by means of mass spectrometry. Analyzing a characteristic of each droplet may include heating each droplet, or

Claim 22 and 29 novelty under PCT Article 33(2) as being anticipated by Anderson et al. US 6,620,620.

surface onto which each droplet is deposited may be applied to assist in the formation of atomized spray. See [0009]

Anderson et al. disclose in FIGS. 2A and 2B illustrate the arrangements used in the droplet sensing method known as video drop sensing. Droplet 42 is deposited from needle 36 of syringe 34 onto deposition surface plate 44. Light source 66 directs light through light diffuser 68, droplet 42 and optional diffuser 70 to video camera 72. As shown in FIG. 2B, video screen 74 (which may be a component of display 24) includes crosshairs 76 which are movable by controls 26 so as to permit the operator to determine a change in the light obscuration as the droplet evaporates. Video camera 72 is interfaced with the power, logic and computing module 22 shown in FIG. 1 so that a change in light obscuration due to the evaporation of at least a portion of the liquid or solvent permits the next successive droplet to be deposited in a timely manner. In a preferred embodiment, a blue or other colored filter may be used to improve the sensitivity of the video camera, and a blunt end needle may be used to improve the wetting of the droplet to the needle.

applying a pneumatic or explosive force to each droplet, so as to form an atomized spray and determining a characteristic each droplet by means of mass spectrometry. Each droplet may be vibrated so as to cause atomization, whereupon a characteristic of each droplet can be determined by means of mass spectrometry. Vibrating the droplet may include focusing a pulsed laser (light source) onto the surface or backside of the surface in a proximity of each droplet, utilizing acoustic waves, or mechanically vibrating the surface. A voltage to the

International application No. PCT/US05/02033

Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

FIG. 3 is a visual representation of the sensing of the evaporative process by means of the change in a reflected laser beam. Droplet 42 is deposited from needle 36 of syringe 34 onto deposition surface plate 44. Laser 78 directs a laser beam at the droplet on deposition surface plate 44, which beam is reflected toward detector 80. When the liquid or solvent droplet has not yet fully evaporated, the reflected beam is scattered and the detector 80 does not yet receive the full beam. Detector 80 is interfaced with the power, logic, and computing module 22 shown in FIG. 1 so that a change in the spectral distribution of the reflected laser beam due to the evaporation of at least a portion of the liquid or solvent permits the next successive droplet to be deposited in a timely manner.

Claims 1, 3, 10, 12, 22, 29 and 31 novelty under PCT Article 33(2) as being anticipated by Yuguchi et al. US 5,275,787.

Yuguchi et al. discloses In FIGS. 9 and 10, a fixed transparent glass plate 16 is arranged tilted relative to the discharge axis 1 of the nozzle 1. The laser beam emitted from the laser light source 7 is incident from the back of the glass plate 16. The discharge axis 1 of the nozzle 1 and the optical axis O of the laser light are arranged so as to cross near the glass plate 16. The detector 5 is arranged on the optical axis of the laser light. A condenser lens (not shown) and a beam stopper (not shown) for preventing the laser beam from directly entering the detector 5 are provided in front of the detector 5 on the optical axis. The condenser lens and beam stopper form a dark-field optical system so that light scattered in forward directions of the optical axis by a particle S situated at a measuring position on the glass plate 16 onto which the laser beam is projected is subjected to photometry by the detector 5. The detector 6 is arranged in a direction crossing each of the optical axis O of the laser light and the axis 1 of the liquid drop discharge. A condenser lens (not shown) and the wavelength selection filter 15 are provided in front of the detector 6 so that fluorescence emitted from the particle S at the measuring position is selectively subjected to photometry by the detector 6.

The size of the opening and the capacity of the heater are set so that the liquid drops discharged from the nozzle have diameters of about $50 \mu m-80 \mu m$.

Claims 22 and 29 novelty under PCT Article 33(2) as being anticipated by Krause et al. US 5,586,200.

Krause et al. teach a system for measuring sample volumes of droplets using a rod-like transfer element. The light of lamp (1) is split into two beams (2) and (3) with each beam being directed onto the sample adhering to the transfer element (4) via a system of lenses, diaphragms, and mirrors. The axis of a rod-like transfer element runs perpendicularly to the drawing surface. The light beams (2) and (3) consequently illuminate the sample droplets from two perpendicular directions which in turn run perpendicularly to the axis of the transfer element. The illuminated sample droplet is located in the focus of the lens systems (5) and (6) by which it is imaged in such a way that a sharp picture is generated on the CCD-camera (8). The bundles of beams emerging from the lens systems (5) and (6) pass through an arrangement of mirrors, diaphragms, and lenses. The semi-permeable mirror (7) directs the partial beams onto the CCD-camera (8).

Claims 4-9, 11, 13-21, 23-28, 30, and 32-36 meet the criteria set out in PCT Article 33(2)-(3), because the prior art does not teach or fairly suggest and immiscible organic liquid that controls evportation, a second immiscible liquid, a thermal gradient in the immiscible liquid, and a dye droplet in the immiscible liquid.

Claims 1-36 meet the criteria set out in PCT Article 33(4), and thus meet industrial applicability because the subject matter claimed can be made or used in industry.